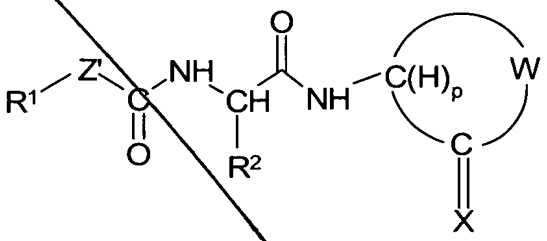
amino, aminoacyl, acylamino, alkaryl, aryl, aryloxy, azido, carboxyl, carboxylalkyl, cyano, halo, nitro, heteroaryl, heterocyclic, aminoacyloxy, oxyacylamino, thioalkoxy, substituted thioalkoxy, thioaryloxy, thioheteroaryloxy, -SO-alkyl, -SO-substituted alkyl, -SO-aryl, -SO-heteroaryl, -SO_2-alkyl, -SO_2-substituted alkyl, -SO_2-aryl, -SO_2-heteroaryl, trihalomethyl, mono- and di-alkylamino, mono- and di-(substituted alkyl)amino, mono- and di-arylamino, mono- and di-heteroarylamino, mono- and di-heterocyclic amino, and unsymmetric di-substituted amines having different substituents selected from alkyl, substituted alkyl, aryl, heteroaryl and heterocyclic, and the like. Preferred substituents include alkyl, alkoxy, halo, cyano, nitro, trihalomethyl, and thioalkoxy.--

IN THE CLAIMS

Please replace Claims 91, 93, 118 and 120 with the following:

91. A pharmaceutical composition comprising a pharmaceutically inert carrier and a pharmaceutically effective amount of formula IA:



IA

wherein R^1 is selected from the group consisting of:

A) alkyl of from 1 to 10 carbon atoms;

B) alkenyl of from 2 to 10 carbon atoms and 1-2 sites of alkenyl unsaturation;

C) alkynyl of from 2 to 10 carbon atoms and from 1-2 sites of alkynyl unsaturation;

D) cycloalkyl of from 3 to 12 carbon atoms;

E) cycloalkenyl of from 4 to 8 carbon atoms;